

AMENDMENTS TO THE CLAIMS

1. **(Currently amended)** A method for potentiating a morphogen activity in a neuron, comprising administering to a mammal contacting the neuron with a composition, the composition comprising a molecule which: that
 - (a) is a neutropoietic cytokine antagonist, a retinoid antagonist, or a cAMP-dependent messenger pathway inhibitor; and
 - (b) overcomes morphogen inhibition of the morphogen activity *in vitro*; thereby potentiating the morphogen activity in a neuron.
2. **(Currently amended)** A method for promoting neuronal cell growth, comprising administering to a mammal contacting a neuron with a composition, the composition comprising a molecule which: that
 - (a) is a neutropoietic cytokine antagonist, a retinoid antagonist, or cAMP-dependent messenger pathway inhibitor; and
 - (b) overcomes morphogen inhibition, of so as to potentiate growth-promoting effects of endogenous morphogens *in vitro*; thereby promoting neuronal cell growth.
- 3 - 4. **(Canceled)**
5. **(Previously presented)** The method of claim 1, wherein said morphogen activity is endogenous.
6. **(Previously presented)** The method of claim 1, wherein said morphogen activity is the result of an exogenously provided morphogen.
7. **(Currently amended)** The method of claim 42, wherein said composition further comprises a morphogen.
8. **(Currently amended)** The method of claim 3 or 41 or 2, wherein said disorder neuron is injured by Alzheimer's disease, Parkinson's disease, Huntington's disease, senile dementia, alcohol-induced dementia, or stroke.

9 -10. **(Canceled)**

11. **(Currently amended)** The method of claim 10-1 or 2, wherein said ~~neuropoetic~~ ~~neuropoietic~~ cytokine antagonist is an LIF (Leukemia-Inhibitory Factor) antagonist or a CNTF (Ciliary Neurotrophic Factor) antagonist.
12. **(Currently amended)** The method of claim 11, wherein said ~~neuropoetic cytokine antagonist is a~~ LIF (Leukemia-Inhibitory Factor) antagonist ~~which~~ is a monoclonal antibody to a gp130 protein.

13 – 15. **(Canceled)**

16. **(Previously presented)** The method of claim 7, wherein said morphogen comprises an amino acid sequence selected from a sequence:
 - (a) having at least 70% homology with the C-terminal seven-cysteine skeleton of human OP-1 (Osteogenic Protein 1), residues 330-431 of SEQ ID NO: 2;
 - (b) having greater than 60% amino acid sequence identity with said C-terminal seven-cysteine skeleton of human OP-1;
 - (c) defined by Generic Sequence 7, SEQ ID NO: 4;
 - (d) defined by Generic Sequence 8, SEQ ID NO: 5;
 - (e) defined by Generic Sequence 9, SEQ ID NO: 6;
 - (f) defined by Generic Sequence 10, SEQ ID NO: 7; or
 - (g) defined by OPX, SEQ ID NO: 3.

17. **(Previously presented)** The method of claim 7, wherein said morphogen is human OP-1 (Osteogenic Protein 1), mouse OP-1, human OP-2 (Osteogenic Protein 2), mouse OP-2, 60A, GDF-1 (Growth/Differentiation Factor-1), BMP2A (Bone Morphogenesis Protein 2A), BMP2B (Bone Morphogenesis Protein 2B), DPP (Decapentaplegic), Vgl, Vgr-1 (Vg1-related sequence), BMP3 (Bone Morphogenesis Protein 3), BMP5 (Bone Morphogenesis Protein 5), or BMP6 (Bone Morphogenesis Protein 6).
18. **(Previously presented)** The method of claim 7, wherein said morphogen is OP-1.

19. **(Currently amended)** The method of claim 1, wherein the molecule neuropoietic cytokine antagonist binds an endogenous ligand for a cytokine receptor-or a retinoid receptor.

20 – 21. **(Canceled)**

22. **(Currently amended)** The method of claim 191 or 2, wherein the molecule which binds an endogenous ligand for a retinoid receptor retinoid antagonist is a retinoic acid receptor or retinoid X receptor.

23 - 24. **(Canceled)**

25. **(Currently amended)** The method of claim 241, wherein said cAMP-dependent messenger pathway inhibitor comprises a protein kinase A inhibitor.

26. **(Previously presented)** The method of claim 25, wherein said protein kinase A inhibitor is (2-p-bromocinnamylaminoethyl)-5-isoquinolinesulfonamide, an enantiomer of dibutyryl cAMP, or an enantiomer of cAMP.

27 – 32. **(Canceled)**

33. **(New)** The method of claim 1, wherein the retinoid antagonist binds an endogenous ligand for a retinoid receptor.

34. **(New)** The method of claim 1, wherein said morphogen activity is activity to stimulate dendritic growth.

35. **(New)** The method of claim 1, wherein said morphogen activity is activity of OP-1.

36. **(New)** The method of claim 2, wherein said neuronal cell growth is dendritic growth.

37. **(New)** The method of claim 1, 2, 34 or 36, wherein said neuron is a sympathetic neuron.

38. **(New)** The method of claim 11, wherein said CNTF inhibitor is phosphatidylinositol-specific phospholipase C.